

USE OF 2 α -METHYL-19-NOR-20(S)-1 α ,25-DIHYDROXYVITAMIN D₃
TO INCREASE BONE STRENGTH

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Attorney Docket No.: 1256-00922
Client Reference No.: P _____

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BACKGROUND OF THE INVENTION

5 This invention relates to vitamin D compounds, and more particularly to pharmaceutical uses for 2 α -methyl-19-nor-20(S)-1 α ,25-dihydroxyvitamin D₃.

The natural hormone, 1 α ,25-dihydroxyvitamin D₃ and its analog in ergocalciferol series, i.e. 1 α ,25-dihydroxyvitamin D₂ are known to be highly potent regulators of calcium homeostasis in animals and humans, and more recently their activity in cellular differentiation has been established, Ostrem et al., Proc. Natl. Acad. Sci. USA, 84, 2610 (1987). Many structural analogs of these metabolites have been prepared and tested, including 1 α -hydroxyvitamin D₃, 1 α -hydroxyvitamin D₂, various side chain homologated vitamins and fluorinated analogs. Some of these compounds exhibit an interesting separation of activities in cell differentiation and calcium regulation. This difference in activity may be useful in the treatment of a variety of diseases as renal osteodystrophy, vitamin D-resistant rickets, osteoporosis, psoriasis, and certain malignancies.

15 Recently, a new class of vitamin D analogs has been discovered, i.e. the so called 19-nor-vitamin D compounds, which are characterized by the replacement of the A-ring exocyclic methylene group (carbon 19), typical of the vitamin D system, by two hydrogen atoms. Biological testing of such 19-nor-analogs (e.g., 1 α ,25-dihydroxy-19-nor-vitamin D₃) revealed a selective activity profile with high potency in inducing cellular differentiation, and very low calcium mobilizing activity. Thus, these compounds are potentially useful as therapeutic agents for the treatment of malignancies, or the treatment of various skin disorders. Two different methods of synthesis of such 19-nor-vitamin D analogs have been described (Perlman et al., Tetrahedron Lett. 31, 1823 (1990); Perlman

et al., Tetrahedron Lett. 32, 7663 (1991), and DeLuca et al., U.S. Pat. No. 5,086,191).

In U.S. Pat. No. 4,666,634, 2 β -hydroxy and alkoxy (e.g., ED-71) analogs of 1 α ,25-dihydroxyvitamin D₃ have been described and examined by Chugai group as potential drugs for osteoporosis and as antitumor agents. See also
5 Okano et al., Biochem. Biophys. Res. Commun. 163, 1444 (1989). Other 2-substituted (with hydroxyalkyl, e.g., ED-120, and fluoroalkyl groups) A-ring analogs of 1 α ,25-dihydroxyvitamin D₃ have also been prepared and tested (Miyamoto et al., Chem. Pharm. Bull. 41, 1111 (1993); Nishii et al., Osteoporosis
10 Int. Suppl. 1, 190 (1993); Posner et al., J. Org. Chem. 59, 7855 (1994), and J. Org. Chem. 60, 4617 (1995)).

Recently, 2-substituted analogs of 1 α ,25-dihydroxy-19-nor-vitamin D₃ have also been synthesized, i.e. compounds substituted at 2-position with hydroxy or alkoxy groups (DeLuca et al., U.S. Pat. No. 5,536,713), which exhibit interesting
15 and selective activity profiles. All these studies indicate that binding sites in vitamin D receptors can accommodate different substituents at C-2 in the synthesized vitamin D analogs.

In a continuing effort to explore the 19-nor class of pharmacologically important vitamin D compounds, an analog which is characterized by the presence of a methyl substituent at the carbon 2 (C-2) has been synthesized and
20 tested. Of particular interest is the analog which is characterized by the unnatural configuration of the methyl group at carbon 20 (C-20), i.e. 2 α -methyl-19-nor-20(S)-1 α ,25-dihydroxyvitamin D₃. This vitamin D analog seemed an interesting target because the relatively small methyl group at C-2 should not interfere with
25 the vitamin D receptor. Moreover, molecular mechanics studies performed on the model 1 α -hydroxy-2-methyl-19-nor-vitamins indicate that such molecular modification does not change substantially the conformation of the cyclohexanediol ring A. However, introduction of the 2-methyl group into 19-nor-

vitamin D carbon skeleton changes the character of its 1α - and 3β - A-ring hydroxyls. Both hydroxyls are allylic to the exocyclic methylene group similar to the 1α -hydroxyl group (crucial for biological activity) in the molecule of the natural hormone, $1\alpha,25-(\text{OH})_2\text{D}_3$.

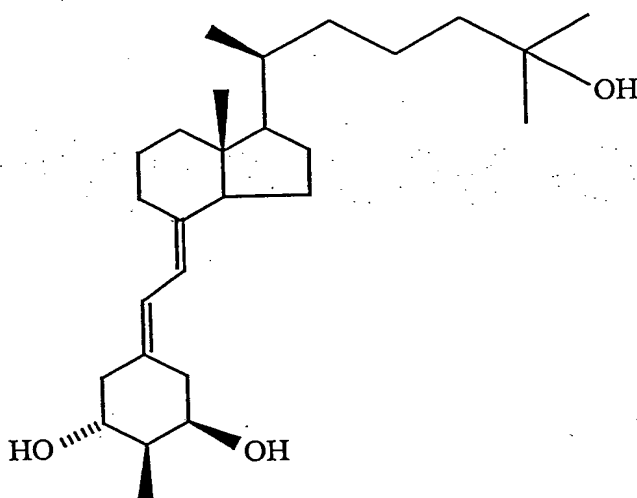
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SUMMARY OF THE INVENTION

The present invention is directed toward 2α -methyl-19-nor-20(S)- $1\alpha,25$ -dihydroxyvitamin D_3 , its biological activity, and various pharmaceutical uses for this compound.

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Structurally this 19-nor analog is characterized by the general formula I shown below:



The solid wedge-shaped line to the methyl substituent at C-20 indicates that carbon 20 has the S configuration.

The above compound exhibits a desired, and highly advantageous, pattern of biological activity. This compound is characterized by intestinal calcium transport activity equal to that of $1\alpha,25$ -dihydroxyvitamin D_3 , but exhibiting relatively high activity, as compared to $1\alpha,25$ -dihydroxyvitamin D_3 , in its ability to mobilize calcium from bone. Hence, this compound is highly specific in its calcemic activity. Its preferential activity on mobilizing calcium from bone allows

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the in vivo administration of this compound for the treatment of metabolic bone diseases where bone loss is a major concern. Because of its preferential activity on bone, this compound would be a preferred therapeutic agent for the treatment of diseases where bone formation is desired, such as osteoporosis, especially low bone turnover osteoporosis, steroid induced osteoporosis, senile osteoporosis or postmenopausal osteoporosis, as well as osteomalacia and renal osteodystrophy. The treatment may be transdermal, oral or parenteral. The compound may be present in a composition in an amount from about 0.1 μ g/gm to about 50 μ g/gm of the composition, and may be administered in dosages of from about 0.1 μ g/day to about 10 μ g/day.

The compound of the invention is also especially suited for treatment and prophylaxis of human disorders which are characterized by an imbalance in the immune system, e.g. in autoimmune diseases, including multiple sclerosis, diabetes mellitus, host versus graft reaction, and rejection of transplants; and additionally for the treatment of inflammatory diseases, such as rheumatoid arthritis and asthma, as well as the improvement of bone fracture healing and improved bone grafts. Acne, alopecia, skin conditions such as dry skin (lack of dermal hydration), undue skin slackness (insufficient skin firmness), insufficient sebum secretion and wrinkles, and hypertension are other conditions which may be treated with the compound of the invention.

The above compound is also characterized by high cell differentiation activity. Thus, this compound also provides a therapeutic agent for the treatment of psoriasis, or as an anti-cancer agent, especially against leukemia, colon cancer, breast cancer and prostate cancer. The compound may be present in a composition to treat psoriasis in an amount from about 0.01 μ g/gm to about 50 μ g/gm of the composition, and may be administered topically, transdermally, orally or parenterally in dosages of from about 0.01 μ g/day to about 10 μ g/day.

It has also been discovered that this compound increases breaking strength (cortical strength) as well as crushing strength (trabecular strength) of bones. Thus, this compound could also be used in conjunction with bone replacement procedures such as hip replacements, knee replacements, and the like.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a graph illustrating the relative activity of 2α -methyl-19-nor-20(S)- $1\alpha,25$ -dihydroxyvitamin D_3 , 2α -methyl-19-nor- $1\alpha,25$ -dihydroxyvitamin D_3 and $1\alpha,25$ -dihydroxyvitamin D_3 to compete for binding of $[^3H]$ - $1,25$ -(OH) $_2$ - D_3 to the vitamin D pig intestinal nuclear receptor;

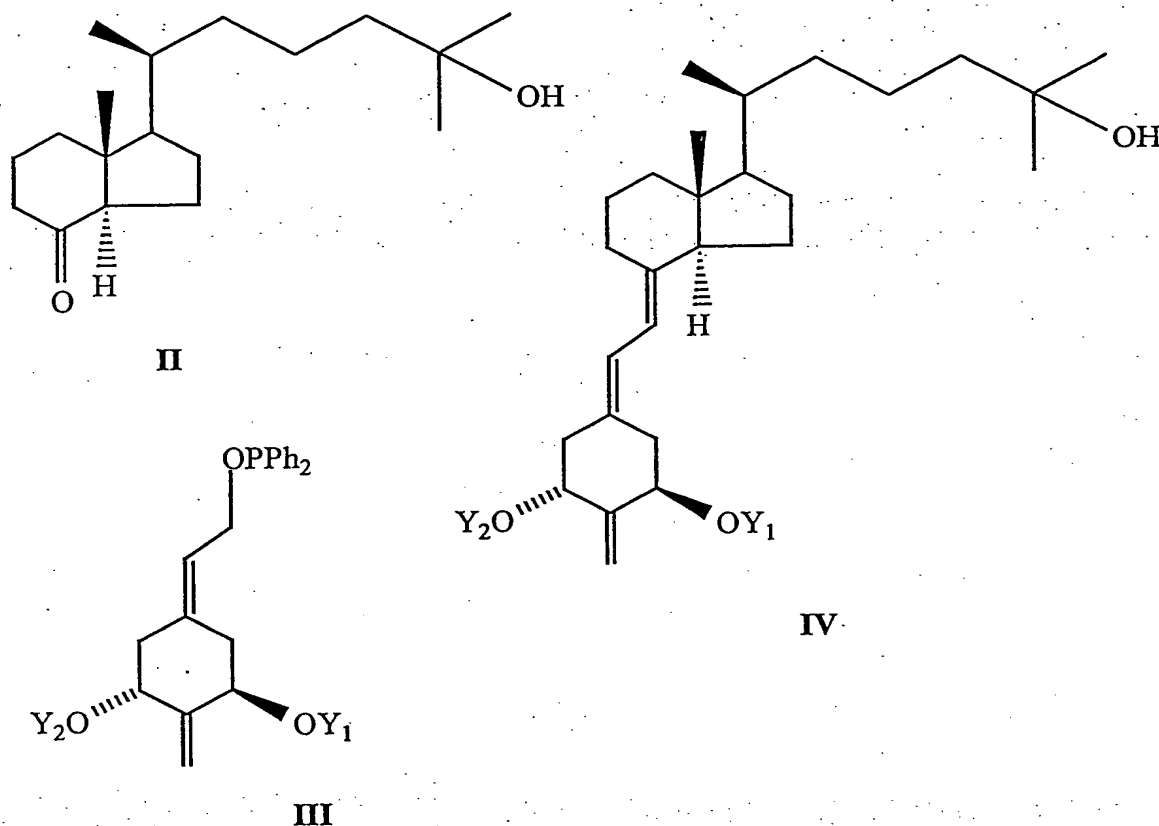
Figure 2 is a graph illustrating the bone mineral density in ovariectomized old female rats as a result of treatment with 2α -methyl-19-nor-20(S)- $1\alpha,25$ -dihydroxyvitamin D_3 as compared to $1\alpha,25$ -dihydroxyvitamin D_3 ; and

Figure 3 is a graph illustrating the percent HL-60 cell differentiation as a function of the concentration of 2α -methyl-19-nor-20(S)- $1\alpha,25$ -dihydroxyvitamin D_3 , 2α -methyl-19-nor- $1\alpha,25$ -dihydroxyvitamin D_3 and $1\alpha,25$ -dihydroxyvitamin D_3 .

DETAILED DESCRIPTION OF THE INVENTION

2α -methyl-19-nor-20(S)- $1\alpha,25$ -dihydroxyvitamin D_3 (referred to herein as 2AMD) was synthesized and tested. Structurally, this 19-nor analog is characterized by the general formula I previously illustrated herein.

The preparation of 2α -methyl-19-nor-20(S)- $1\alpha,25$ -dihydroxyvitamin D_3 having the basic structure I can be accomplished by a common general method, i.e. the condensation of a bicyclic Windaus-Grundmann type ketone II with the allylic phosphine oxide III to the corresponding 2-methylene-19-nor-vitamin D analog IV followed by selective reduction of the exomethylene group at C-2 and deprotection at C-1 and C-3 in the latter compound:



In the structures II, III, and IV groups Y_1 and Y_2 are hydroxy-protecting groups, it being also understood that any functionalities that might be sensitive, or that
 5 interfere with the condensation reaction, be suitably protected as is well-known in the art. The process shown above represents an application of the convergent synthesis concept, which has been applied effectively for the preparation of vitamin D compounds [e.g. Lythgoe et al., J. Chem. Soc. Perkin Trans. I, 590 (1978); Lythgoe, Chem. Soc. Rev. 9, 449 (1983); Toh et al., J. Org. Chem. 48,
 10 1414 (1983); Baggiolini et al., J. Org. Chem. 51, 3098 (1986); Sardina et al., J. Org. Chem. 51, 1264 (1986); J. Org. Chem. 51, 1269 (1986); DeLuca et al., U.S. Pat. No. 5,086,191; DeLuca et al., U.S. Pat. No. 5,536,713].

Hydrindanones of the general structure II are known, or can be prepared by known methods.

For the preparation of the required phosphine oxides of general structure III, a new synthetic route has been developed starting from a methyl quinate derivative which is easily obtained from commercial (1R,3R,4S,5R)-(-)-quinic acid as described by Perlman et al., Tetrahedron Lett. 32, 7663 (1991) and DeLuca et al., U.S. Pat. No. 5,086,191.

The overall process of the synthesis of compound I is illustrated and described more completely in U.S. Patent No. 5,945,410 issued August 31, 1999 and entitled "2-Alkyl-19-Nor-Vitamin D Compounds" the specification of which is specifically incorporated herein by reference.

BIOLOGICAL ACTIVITY OF 2 α -METHYL-20(S)- 19-NOR-1,25-(OH)₂D₃

The introduction of a methyl group in the 2 α -position of the 20(S) isomer of 19-nor-1,25-(OH)₂D₃ had little or no effect on binding to the porcine intestinal vitamin D receptor. This compound bound equally well to the porcine receptor as compared to the standard 1,25-(OH)₂D₃ (Figure 1). It might be expected from these results that this compound would have equivalent biological activity. Surprisingly, however, the 2 α -methyl and 20(S) substitutions produced a highly selective analog with its primary action on bone.

Figure 2 shows that 2AMD is extraordinarily effective in building bone mass in ovariectomized rats as compared to the native hormone without increasing serum calcium concentration. This is as yet an unprecedented new finding for a vitamin D compound.

Figure 3 illustrates that 2AMD is 50-100 times more potent than 1,25(OH)₂D₃ on HL-60 differentiation, making it an excellent candidate for the treatment of psoriasis and cancer, especially against leukemia, colon cancer, breast cancer and prostate cancer.

increased bone mass provided by 2AMD translates into marked increases in bone strength. This increased strength to fracture in femur shows cortical strength while increased strength to crush fractures of vertebra illustrates trabecular bone strength (Table 2). Interestingly, even the percent ash is unexpectedly increased further by 2AMD. Of great importance is that at the dosage levels used in this study, there was no change in serum calcium of animals that showed the marked elevation of bone mass. This argues that a window of safety exists between the use of 2AMD to increase bone mineral content and the action of 2AMD in elevating serum calcium.

When given for 7 days in a chronic mode, the most potent individual compound tested was 2α -methyl 19-nor-20S-1,25-(OH) $_2$ D $_3$ (Table 3). When given at 130 pmol/day, the activity of this compound on bone calcium mobilization (serum calcium) was much higher than that of the native hormone, possible as high as 10 or 100 times higher. Under identical conditions, twice the dose of 1,25-(OH) $_2$ D $_3$ gave a serum calcium value of 6.6 ± 0.4 mg/100 ml, while 2α -methyl-19-nor-20S-1,25-(OH) $_2$ D $_3$ gave a value of 8.3 ± 0.7 mg/100 ml of serum calcium at the 130 pmol dose. When given at 260 pmol/day, 2α -methyl-19-nor-20S-1,25-(OH) $_2$ D $_3$ produced the astounding value of 10.3 ± 0.11 mg/100 ml of serum calcium at the expense of bone. To show its selectivity, this compound also produced a significant change in intestinal calcium transport at both the 260 pmol and the 130 pmol dose levels while having a strong bone calcium mobilizing activity. At the higher dose, the 2α -methyl-20(S) compound did produce a significant intestinal transport response but also gave an enormous bone mobilization response. With respect to the 2β -methyl-19-nor-20(S) compound, the data in Table 3 show it has little, if any, intestinal calcium transport activity, and little, if any, bone mobilization activity. Thus, the 2α -methyl-19-nor-20(S)-derivative showed strong preferential bone calcium mobilizing activity. These

results illustrate that the 20(S)-2 α -methyl derivative of 19-nor-1,25-(OH)₂D₃ is selective for mobilization of calcium from bone.

These results illustrate that 2AMD is an excellent candidate for an anti-osteoporosis therapy and that it may be useful in a number of other
5 circumstances such as autoimmune diseases, cancer, and psoriasis.

Table 1

Treatment of Ovariectomized Rats with 1,25-(OH)₂D₃ and 2AMD

Group	Treatment	Treatment Time (Weeks)	BMD (g/cm ²)	BMC (g)	Body Wt. (g)	BMC/Body Wt. (mg/g)	Serum CA (mg/dl)	Femur Ash (%)	Femur Ash (mg)
OVX Control	Oil Vehicle/5X/Week	8	0.294±0.004	8.64±3.30	414±15	21.4±1.20	---	---	---
		17	0.296±0.003	9.34±0.50	422±19	22.3±1.69	---	---	---
		30	0.296±0.003	9.41±0.45	404±24	23.4±1.60	11.1±0.17	59.2±0.82	386±21.6
Sham Operated	Oil Vehicle/5X/Week	8	0.302±0.003	9.34±0.38	356±14	26.3±0.76	---	---	---
		17	0.300±0.002	9.14±0.54	351±15	26.4±0.82	---	---	---
		30	0.297±0.004	9.20±0.53	340±13	26.7±1.20	11.8±0.20	81.5±1.20	400±18.0
1,25(OH) ₂ D ₃	250 pmol/d/5X/Week	8	0.297±0.001	8.90±0.40	399±9.3	22.4±0.48	---	---	---
		17	0.308±0.008	9.6±0.39	394±11	24.5±0.87	11.4±0.21	60.8±1.1	417±23
		30	0.310±0.007	10.1±0.30	392±16	26.1±0.97	---	---	---
1,25(OH) ₂ D ₃	500 pmol/d/5X/Week 3X/Week 3X/Week	8	0.312±0.005	10.2±0.40	397±14.2	26.3±0.57	---	---	---
		17	0.331±0.008	11.5±0.25	421±12.8	27.6±0.68	11.9±0.20	61.4±1.3	478±7.5
		30	0.328±0.003	11.8±0.23	432±23.0	28.0±0.69	---	---	---
2AMD	32 pmol/d/2X/Week	8	0.297±0.002	11.2±0.55	371±13.5	31.0±2.50	---	---	---
		17	0.308±0.001	9.7±0.24	401±9.6	24.2±1.04	10.4±0.53	61.7±1.2	447±23.3
		30	0.320±0.003	11.2±0.55	371±13.5	31.0±2.50	---	---	---
2AMD	65 pmol/d/1X/Week	8	0.298±0.002	9.1±0.31	384±9.7	23.9±0.37	---	---	---
		17	0.308±0.004	9.5±0.57	396±12.8	24.3±0.53	10.8±0.22	60.0±1.4	403±12.
		30	0.308±0.003	9.7±0.28	387±13.7	25.1±1.00	---	---	---

All animals were ovariectomized except the sham-operated controls. Values are expressed as mean ± SEM.

Table 2

Strength of Femurs and Vertebrae to Mechanical Stress

Group	Treatment	Stress Value Femur	Stress Value Vertebra
OVX Control	Oil Vehicle/5X/Week	109.31±19.60	14.26±3.58
Sham-Operated	Oil Vehicle/5X/Week	121.36±12.5	13.67±1.79
1,25(OH) ₂ D ₃	250 pmol/day/5X/Week	118.21±19.85	19.24±5.66
1,25(OH) ₂ D ₃	500 pmol/d/3-5X/Week	116.47±16.20	17.14±0.52
2AMD	32 pmol/d/2X/Week	132.19±19.69	20.22±8.53
2AMD	65 pmol/d/1X/Week	114.08±21.71	17.13±4.67

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Table 3

Response of Intestinal Calcium Transport and
Serum Calcium (Bone Calcium Mobilization) Activity
to Chronic Doses of the 20(S) Isomers of 2-Methyl
Derivatives of 19-Nor-1,25-(OH)₂D₃

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Group	Dose (pmol/day/7 days)	Intestinal Calcium Transport (S/M)	Serum Calcium (mg/100 ml)
Vitamin D Deficient	Vehicle	2.9±0.2	4.2±0.1
1,25-(OH) ₂ D ₃ Treated	260	4.6±0.2	6.6±0.4
2α-Methyl-19-Nor-20(S)-1,25-(OH) ₂ D ₃	130	12.9±1.9	8.3±0.7
	260	8.4±1.1	10.3±0.11
2β-Methyl-19-nor-20(S)-1,25-(OH) ₂ D ₃	130	2.9±0.3	4.4±0.1
	260	3.8±0.1	4.4±0.1

With respect to the data in Table 3, male weanling rats were obtained from Sprague Dawley Co. (Indianapolis, IN) and fed a 0.47% calcium, 0.3% phosphorus vitamin D-deficient diet for 1 week and then given the same diet containing 0.02% calcium, 0.3% phosphorus for 2 weeks. During the last week they were given the indicated dose of compound by intraperitoneal injection in 0.1 ml 95% propylene glycol and 5% ethanol each day for 7 days. The control

animals received only the 0.1 ml of 95% propylene glycol, 5% ethanol. Twenty-four hours after the last dose, the rats were sacrificed and intestinal calcium transport was determined everted sac technique as previously described and serum calcium determined by atomic absorption spectrometry on a model 3110
5 Perkin Elmer instrument (Norwalk, CT). There were 5 rats per group and the values represent mean \pm SEM.

For treatment purposes, the compound of this invention defined by formula I may be formulated for pharmaceutical applications as a solution in innocuous solvents, or as an emulsion, suspension or dispersion in suitable
10 solvents or carriers, or as pills, tablets or capsules, together with solid carriers, according to conventional methods known in the art. Any such formulations may also contain other pharmaceutically-acceptable and non-toxic excipients such as stabilizers, anti-oxidants, binders, coloring agents or emulsifying or taste-modifying agents.

15 The compound may be administered orally, topically, parenterally or transdermally. The compound is advantageously administered by injection or by intravenous infusion or suitable sterile solutions, or in the form of liquid or solid doses via the alimentary canal, or in the form of creams, ointments, patches, or similar vehicles suitable for transdermal applications. Doses of from 0.1 μ g to
20 10 μ g per day of the compounds are appropriate for treatment purposes, such doses being adjusted according to the disease to be treated, its severity and the response of the subject as is well understood in the art. Since the compound exhibits specificity of action, each may be suitably administered alone, or together with graded doses of another active vitamin D compound -- e.g. 1 α -
25 hydroxyvitamin D₂ or D₃, or 1 α ,25-dihydroxyvitamin D₃ -- in situations where different degrees of bone mineral mobilization and calcium transport stimulation is found to be advantageous.

Compositions for use in the above-mentioned treatment of psoriasis and other malignancies comprise an effective amount of the 2 α -methyl-20(S)-19-nor-vitamin D compound as defined by the above formula I as the active ingredient, and a suitable carrier. An effective amount of such compound for use in accordance with this invention is from about 0.01 μ g to about 50 μ g per gm of composition, and may be administered topically, transdermally, orally or parenterally in dosages of from about 0.1 μ g/day to about 10 μ g/day.

The compound may be formulated as creams, lotions, ointments, topical patches, pills, capsules or tablets, or in liquid form as solutions, emulsions, dispersions, or suspensions in pharmaceutically innocuous and acceptable solvent or oils, and such preparations may contain in addition other pharmaceutically innocuous or beneficial components, such as stabilizers, antioxidants, emulsifiers, coloring agents, binders or taste-modifying agents.

The compound is advantageously administered in amounts sufficient to effect the differentiation of promyelocytes to normal macrophages. Dosages as described above are suitable, it being understood that the amounts given are to be adjusted in accordance with the severity of the disease, and the condition and response of the subject as is well understood in the art.

The formulations of the present invention comprise an active ingredient in association with a pharmaceutically acceptable carrier therefore and optionally other therapeutic ingredients. The carrier must be "acceptable" in the sense of being compatible with the other ingredients of the formulations and not deleterious to the recipient thereof.

Formulations of the present invention suitable for oral administration may be in the form of discrete units as capsules, sachets, tablets or lozenges, each containing a predetermined amount of the active ingredient; in the form of a powder or granules; in the form of a solution or a suspension in an aqueous liquid

or non-aqueous liquid; or in the form of an oil-in-water emulsion or a water-in-oil emulsion.

Formulations for rectal administration may be in the form of a suppository incorporating the active ingredient and carrier such as cocoa butter, or in the form of an enema.

Formulations suitable for parenteral administration conveniently comprise a sterile oily or aqueous preparation of the active ingredient which is preferably isotonic with the blood of the recipient.

Formulations suitable for topical administration include liquid or semi-liquid preparations such as liniments, lotions, applicants, oil-in-water or water-in-oil emulsions such as creams, ointments or pastes; or solutions or suspensions such as drops; or as sprays.

For asthma treatment, inhalation of powder, self-propelling or spray formulations, dispensed with a spray can, a nebulizer or an atomizer can be used. The formulations, when dispensed, preferably have a particle size in the range of 10 to 100 μ .

The formulations may conveniently be presented in dosage unit form and may be prepared by any of the methods well known in the art of pharmacy. By the term "dosage unit" is meant a unitary, i.e. a single dose which is capable of being administered to a patient as a physically and chemically stable unit dose comprising either the active ingredient as such or a mixture of it with solid or liquid pharmaceutical diluents or carriers.